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plants and lower animals that cause cell agglutination in the presence of certain antigens. These substances are used to detect blood group antigens for in vitro diagnostic purposes.

(b) Classification. Class II (special controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §864.9.

[45 FR 60648, Sept. 12, 1980, as amended at 63 FR 59226, Nov. 3, 1998]

# §864.9575 Environmental chamber for storage of platelet concentrate.

- (a) *Identification*. An environmental chamber for storage of platelet concentrate is a device used to hold platelet-rich plasma within a preselected temperature range.
- (b) Classification. Class II (special controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §864.9.

[45 FR 60648, Sept. 12, 1980, as amended at 63 FR 59226, Nov. 3, 1998]

# §864.9600 Potentiating media for in vitro diagnostic use.

- (a) *Identification*. Potentiating media for in vitro diagnostic use are media, such as bovine albumin, that are used to suspend red cells and to enhance cell reactions for antigen-antibody testing.
- (b) Classification. Class II (special controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §864.9.

[45 FR 60649, Sept. 12, 1980, as amended at 63 FR 59226, Nov. 3, 1998]

## §864.9650 Quality control kit for blood banking reagents.

- (a) *Identification*. A quality control kit for blood banking reagents is a device that consists of sera, cells, buffers, and antibodies used to determine the specificity, potency, and reactivity of the cells and reagents used for blood banking.
- (b) Classification. Class II (performance standards).

[45 FR 60649, Sept. 12, 1980]

# §864.9700 Blood storage refrigerator and blood storage freezer.

- (a) *Identification*. A blood storage refrigerator and a blood storage freezer are devices intended for medical purposes that are used to preserve blood and blood products by storing them at cold or freezing temperatures.
- (b) Classification. Class II (special controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §864.9.

[45 FR 60650, Sept. 12, 1980, as amended at 63 FR 59226, Nov. 3, 1998]

#### §864.9750 Heat-sealing device.

- (a) *Identification*. A heat-sealing device is a device intended for medical purposes that uses heat to seal plastic bags containing blood or blood components.
- (b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §864.9.

[45 FR 60650, Sept. 12, 1980, as amended at 65 FR 2311, Jan. 14, 2000]

### § 864.9875 Transfer set.

- (a) *Identification*. A transfer set is a device intended for medical purposes that consists of a piece of tubing with suitable adaptors used to transfer blood or plasma from one container to another.
- (b) Classification. Class II (performance standards).

[45 FR 60651, Sept. 12, 1980]

### Subpart K—Products Used In Establishments That Manufacture Human Cells, Tissues, and Cellular and Tissue-Based Products (HCT/Ps)

# §864.9900 Cord blood processing system and storage container.

(a) *Identification*. A cord blood processing system and storage container is a device intended for use in the processing and the storage of cord blood. This device is a functionally closed

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processing system that includes containers, other soft goods, and a centrifugation system for cord blood concentration, and a final container for the cryopreservation and the storage of a cord blood product.

(b) Classification. Class II (special controls). The special control for this device is FDA's guidance document entitled "Class II Special Controls Guidance Document: Cord Blood Processing System and Storage Container." For the availability of this guidance document, see §864.1(d).

[72 FR 4638, Feb. 1, 2007]

### PART 866—IMMUNOLOGY AND MICROBIOLOGY DEVICES

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Sec.

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866.2160 Coagulase plasma.

866.2170 Automated colony counter.

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866.2480 Quality control kit for culture media

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866.2540 Microbiological incubator.

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866.2580 Gas-generating device.

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866.3020 Adenovirus serological reagents.

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866.3040 Aspergillus spp. serological reagents.

866.3050 Beta-glucan serological assays.

866.3060 Blastomyces dermatitidis serological reagents.

866.3065 Bordetella spp. serological reagents.

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866.3120 Chlamydia serological reagents.

866.3125 Citrobacter spp. serological reagents. 866.3135 Coccidioides immitis serological reagents.

866.3140 Corynebacterium spp. serological reagents.

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866.3175 Cytomegalovirus serological reagents.

866.3200 Echinococcus spp. serological reagents.

866.3205 Echovirus serological reagents. 866.3210 Endotoxin assay.

866.3220 Entamoeba histolytica serological reagents.

866.3225 Enterovirus nucleic acid assay.

866.3235 Epstein-Barr virus serological reagents.

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866.3250 Erysipelothrix rhusiopathiae sero-

logical reagents. 866.3255 Escherichia coli serological reagents.

866.3270 Flavobacterium spp. serological reagents.

866.3280 Francisella tularensis serological reagents.

866.3290 Gonococcal antibody test (GAT).

866.3300 Haemophilus spp. serological reagents

866.3305 Herpes simplex virus serological assavs.

866.3310 Hepatitis A virus (HAV) serological assays.

866.3320 Histoplasma capsulatum serological reagents.